

## **PUMP SYSTEMS INCLUDING INJECTABLE GABAPENTIN COMPOSITIONS**

### **RELATED APPLICATIONS**

- [1] This application is a continuation in part application of Serial No. 10/611,459, entitled “A method for treating severe tinnitus”, filed July 1, 2003. This application claims priority to the above-referenced application and also claims priority to Provisional Application Serial No. 60/513682, entitled “INJECTABLE GABAPENTIN COMPOSITIONS”, filed October 23, 2003, and Provisional Application Serial No. 60/513681, entitled “INTRATHECAL GABAPENTIN FOR TREATMENT OF PAIN AND EPILEPSY”, filed on October 23, 2003. Each of the above-referenced applications is herein incorporated by reference in their entirety.

### **FIELD OF THE INVENTION**

- [2] This application relates to medical devices, particularly to therapeutic pump systems that include injectable gabapentin compositions.

### **BACKGROUND OF THE INVENTION**

- [3] Gabapentin is a pharmacological agent that mimics the effects of GABA ( $\gamma$ -aminobutyric acid), but gabapentin does not appear to bind a GABA receptor (e.g., GABA<sub>A</sub> and GABA<sub>B</sub> receptors) or have an effect on GABA uptake. Gabapentin has been found to interact with the alpha-2-delta ( $\alpha_2\delta$ ) subunit of voltage-gated calcium channels. Many of the pharmacological effects of gabapentin may be due to its interaction with voltage-gated calcium channels. It is believed that gabapentin decreases calcium ion flow into a neuron, rendering the neuron less excitable. Inhibition of presynaptic calcium influx may prevent the release of neurotransmitters. Thus, like GABA, gabapentin can dampen overactive neural circuitry.
- [4] Solid formulations of gabapentin, such as NEURONTIN, are currently available for oral administration. Oral gabapentin has been primarily used to treat epilepsy although it has been used off-label to treat neuropathic pain and has recently received an FDA-approval

for the treatment of one type of neuropathic pain, post herpetic neuralgia. Some gabapentin can access the CNS when administered orally, because gabapentin is transported across the gut and the blood-brain barrier. It is believed that gabapentin is transported across the blood-brain barrier via an active and saturable L-amino acid transporter. Thus, the amount of gabapentin reaching CNS sites of action is limited. Because this transporter is saturable, even if the concentration of gabapentin in the plasma is increased, the amount which crosses the blood-brain barrier will remain constant.

- [5] Solutions of gabapentin have been prepared for direct administration to the CNS in preclinical animal studies. In some studies, such solutions have been administered intrathecally as a single bolus or as multiple boluses. In these studies, the solutions contained gabapentin in varying concentrations, generally from about 1 mg/ml to about 30 mg/mL. One study (Wang and Yaksh, 1997) reported the use of a solution of 100 mg/ml gabapentin for a single 10 µl intrathecal bolus (1000 µg) injection in rats. Wang and Yaksh found that the 1000 µg bolus injection of gabapentin caused significant hind limb motor weakness. Generally a bolus injection of 300 µg gabapentin will cause hind limb motor weakness in rats. As 10 µl bolus injections may be used in rats, solutions of gabapentin at a concentration greater than about 30 mg/ml have been of little practical use.
- [6] Generally, solutions of gabapentin used in preclinical animal trials contain 0.9% saline in an attempt to approximate physiological saline. However, gabapentin is zwitterionic and may contribute significantly to tonicity of a solution, depending on the concentration of gabapentin. Thus, solutions having high gabapentin concentrations have high tonicity. The presence of 0.9% saline in solutions having a high concentration of gabapentin increases the tonicity, thereby making the solutions hypertonic relative to physiological fluids such as cerebrospinal fluid. For example, the 100 mg/ml gabapentin solution in 0.9% saline described by Wang and Yaksch has a tonicity of about 925 mOsm, as compared to a tonicity of about 300 mOsm for physiological fluids. Hypertonic solutions, when administered to a subject, can result in tissue damage due to shrinkage of

cells. In the cerebrospinal fluid, which is generally a poorly mixed tissue compartment, local damage and shrinkage due to hypertonic solutions is a more pressing concern. When administered as a small-volume bolus and flushed with saline or barbotaged with CSF, the risks associated with the administration of a hypertonic fluid are minimal. In contrast, continuous infusion of a low volume of fluid into the subarachnoid space can result in prolonged exposure of the spinal tissues adjacent to the catheter tip. In this case, the risks associated with administration of a nonphysiological hypertonic solution are increased.

#### SUMMARY OF THE INVENTION

- [7] An embodiment of the invention provides a system comprising a pump having a reservoir, a catheter coupled to the pump and adapted for delivering a therapeutic agent to a cerebrospinal fluid of a patient, and an injectable gabapentin composition housed in the reservoir and deliverable through the catheter in an amount effective to treat pain in the patient when administered to the cerebrospinal fluid of the patient. In an embodiment, the injectable composition is an injectable solution, and gabapentin is present in the solution at a concentration greater than about 30 mg/mL, and the solution has a tonicity of less than about 900 mOsm. In an embodiment, the injectable gabapentin composition comprises less than 0.9% (w/v) sodium chloride.
- [8] Various embodiments of the invention provide several advantages. For example, solutions having reduced hypertonicity may result in reduced tissue and cell damage due to injection of the solution. By reducing solvent tonicity, increased concentrations of gabapentin may be placed in injectable solutions without rendering the solutions excessively hypertonic. When used in a pump system designed to deliver a therapeutic agent, compositions having increased concentrations of gabapentin will allow for greater time to elapse, relative to compositions having lower gabapentin concentrations, before the pump requires refilling. Increasing time between refills is particularly important when the pump is an implantable pump.

- [9] These and other advantages of the invention will become evident upon reading the description herein.

#### BRIEF DESCRIPTION OF THE DRAWINGS

- [10] Figure 1 is a diagrammatic illustration of a pump system for delivering a composition comprising a therapeutic agent according to an embodiment of the present invention.
- [11] Figure 2 is a diagrammatic illustration of a catheter implanted in a patient according to an embodiment of the present invention.
- [12] Figure 3 is a diagrammatic illustration of an implanted catheter and pump in accordance with an embodiment of the present invention.
- [13] Figure 4 is a diagrammatic illustration of an implanted catheter and pump in accordance with an embodiment of the present invention.
- [14] Figure 5 is a diagrammatic illustration of a catheter and external pump in accordance with an embodiment of the present invention.
- [15] The drawings are not necessarily to scale.

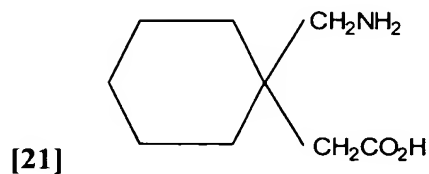
#### DETAILED DESCRIPTION

- [16] The following description illustrates various embodiments of the invention. It is to be understood that other embodiments of the present invention are contemplated and may be made without departing from the scope or spirit of the present invention. Thus, the following description is not to be taken in a limiting sense.
- [17] All scientific and technical terms used in this application have meanings commonly used in the art unless otherwise specified. The definitions provided herein are to facilitate understanding of certain terms used frequently herein and are not meant to limit the scope of the present disclosure.

[18] Embodiments of the present invention provide injectable compositions comprising gabapentin. Injectable compositions comprising gabapentin according to embodiments of the invention may be used for any purpose for which study or use of gabapentin is desired. For example, injectable compositions comprising gabapentin may be used in studies to determine or elucidate (a) the effect of gabapentin on a molecule, cell, tissue, organ, organism, or combination thereof; (b) the mechanism of action of gabapentin, (c) the properties of gabapentin, a solution comprising gabapentin, or a combination thereof; and (d) the like. Injectable compositions comprising gabapentin may also be used as therapy to treat a disease state responsive to gabapentin, such as epilepsy, pain, tinnitus, drug addiction, bipolar disorder, osteoarthritis, migraine, and anxiety disorders including social phobia. In the context of the present invention, the terms "treat", "therapy", and the like are meant to include methods to alleviate, slow the progression, prevent, attenuate, or cure the treated disease.

[19] Injectable Composition

[20] An embodiment of the invention provides an injectable composition comprising gabapentin. As used herein, gabapentin refers to 1-(aminomethyl)cyclohexane acetic acid and pharmaceutically acceptable salts, solvates, hydrates, and polymorphs thereof. 1-(aminomethyl)cyclohexane acetic acid is a  $\gamma$ -aminobutyric acid (GABA) analogue with a molecular formula of  $C_9H_{17}NO_2$  and a molecular weight of 171.24. 1-(aminomethyl)cyclohexane acetic acid is freely soluble in water and in both basic and acidic aqueous solutions. 1-(aminomethyl)cyclohexane acetic acid has a structure of:



[22] Gabapentin may be obtained from a variety of commercial sources, such as Shanghai Zhongxi International Trading Co., Shanghai, China; Hikal Limited, Bangalore, Karnaraka, India; Erregierre S.p.A., San Paolo d'Argon (BG), Italy; MediChem, SA, Sant

Joan Despi (Barcelona), Spain; Ranbaxy Laboratories, New Delhi, India; Procos S.p.A., Cameri, Italy; Zambon Group, Milan, Italy; Hangzhuo Chiral Medicine Chemicals Co., Hangzhuo, China; InterChem Corporation USA, Paramus, NJ; SST Corporation, Clifton, NJ; Teva Pharmaceuticals USA, North Wales, PA; Plantex USA, Hakensack, NJ; and Sigma-Aldrich, St. Louis, MO, or an appropriate distributor. Alternatively, gabapentin may be synthesized and/or prepared as known in the art.

- [23] As used herein, “injectable composition” refers to a composition that is fluid at room temperature, which fluid is capable of being injected into a patient. Injectable compositions include solutions, suspensions, dispersions, and the like. Injectable solutions, suspensions, dispersions, and the like may be formulated according to techniques well-known in the art (see, for example, Remington's Pharmaceutical Sciences, Chapter 43, 14th Ed., Mack Publishing Co., Easton, Pa.), using suitable dispersing or wetting and suspending agents, such as sterile oils, including synthetic mono- or diglycerides, and fatty acids, including oleic acid.
- [24] Injectable compositions comprising gabapentin may be prepared in water, saline, isotonic saline, phosphate-buffered saline, citrate-buffered saline, and the like and may optionally mixed with a nontoxic surfactant. Dispersions may also be prepared in glycerol, liquid polyethylene, glycols, DNA, vegetable oils, triacetin, and the like and mixtures thereof. Under ordinary conditions of storage and use, these preparations may contain a preservative to prevent the growth of microorganisms. Pharmaceutical dosage forms suitable for injection or infusion include sterile, aqueous solutions or dispersions or sterile powders comprising an active ingredient which powders are adapted for the extemporaneous preparation of sterile injectable or infusible solutions or dispersions. Preferably, the ultimate dosage form is a sterile fluid and stable under the conditions of manufacture and storage. A liquid carrier or vehicle of the solution, suspension or dispersion may be a solvent or liquid dispersion medium comprising, for example, water, ethanol, a polyol such as glycerol, propylene glycol, or liquid polyethylene glycols and the like, vegetable oils, nontoxic glyceryl esters, and suitable mixtures thereof. Proper fluidity of solutions, suspensions or dispersions may be maintained, for example, by the

formation of liposomes, by the maintenance of the required particle size, in the case of dispersion, or by the use of nontoxic surfactants. The prevention of the action of microorganisms can be accomplished by various antibacterial and antifungal agents, for example, parabens, chlorobutanol, phenol, sorbic acid, thimerosal, and the like. In many cases, it will be desirable to include isotonic agents, for example, sugars, buffers, or sodium chloride. Prolonged absorption of the injectable compositions can be brought about by the inclusion in the composition of agents delaying absorption--for example, aluminum monostearate hydrogels and gelatin. Excipients that increase solubility, such as cyclodextrin, may be added.

- [25] Sterile injectable compositions may be prepared by incorporating a therapeutic agent in the desired amount in the appropriate solvent with various other ingredients as enumerated above and, as desired, followed by sterilization. Any means for sterilization may be used. For example, the solution may be autoclaved or filter sterilized. In the case of sterile powders for the preparation of sterile injectable solutions, methods of preparation include vacuum drying and freeze-drying techniques, which yield a powder of the active ingredient plus any additional desired ingredient present in a previously sterile-filtered solution.
- [26] Injectable compositions comprising gabapentin may be heat treated or sterilized by autoclaving. Because increased temperature may result in increased conversion of gabapentin to its corresponding lactam, which is generally considered more toxic than gabapentin, it would be expected that high temperatures should be avoided when preparing compositions comprising gabapentin. Surprisingly, compositions comprising gabapentin may be sterilized by autoclaving to provide suitable sterile injectable gabapentin compositions. Heat treatment, whether or not through autoclaving, may be performed at any combination of temperature and time necessary to sterilize a composition comprising gabapentin. For example, a composition may be subjected to heat treatment for about 2 minutes to about 60 minutes at temperatures of about 110 °C to about 140 °C. Specific exemplary times and temperatures that may be used include 24 minutes at 121.1 °C, 4 minutes at 130 °C, 30 min at 118 °C, and 6-8 min at 121.1 °C. It

will be recognized that with higher temperatures and the longer durations of heat treatment, the likelihood of gabapentin lactam formation will be increased. To prevent excess formation of lactam, the time and temperature of heat treatment may be adjusted to a combination that reduces lactam formation, yet continues to sterilize the composition comprising gabapentin.

- [27] In an embodiment, an appropriate weight of non-sterile gabapentin powder is dissolved in an appropriate volume of sterile water for injection to yield an aqueous gabapentin solution. The pH is adjusted to about 6 with 1N NaOH or 1N HCl, and the resulting solution is sterilized by autoclaving.
- [28] In an embodiment, an injectable composition comprising gabapentin is an injectable solution comprising an aqueous solvent. The solvent may be water or saline. The saline may be, *e.g.*, 0.9% (w/v) sodium chloride or a solution where just enough sodium chloride is added to make the final solution isotonic. The saline may be sterile saline. In an embodiment, the final solution has a pH between about 4 and about 9, between about 5 and about 7, between about 5.5 and about 6.5, or about 6. The pH of an injectable gabapentin composition may be adjusted with a pharmacologically acceptable acid, base, buffer or combination thereof. In an embodiment, pH is adjusted with hydrochloric acid or sodium hydroxide. The hydrochloric acid or sodium hydroxide may be in any suitable form, such as a 1N solution. In an embodiment, an injectable gabapentin solution has a pH in the range of between about 4 and about 9, between about 5 and about 7, or about 6. Preferably, the final solution contains less than about 5% of gabapentin lactam. In an embodiment, the final solution contains less than about 2% gabapentin lactam. In an embodiment, the final solution contains less than about 1% gabapentin lactam.
- [29] A composition comprising gabapentin according to an embodiment of the invention includes an amount of gabapentin effective to treat a disease responsive to gabapentin. In an embodiment, the amount of gabapentin is effective to treat a gabapentin-responsive disease when administered intrathecally. Generally, gabapentin may be present in a solution or suspension at a concentration between about 0.1 mg/mL and about 100



mg/mL. In an embodiment, gabapentin is present in a solution or suspension at a concentration between about 10 mg/mL and about 90 mg/mL. In an embodiment, gabapentin is present in a solution or suspension at a concentration between about 20mg/mL and about 80 mg/mL. In an embodiment, gabapentin is present in a solution or suspension at a concentration of about 80 mg/mL.

[30] In an embodiment, an injectable gabapentin composition is substantially free of preservatives, substantially free of buffers, or substantially free of both preservatives and buffers.

[31] Tonicity

[32] In an embodiment, the invention provides an injectable composition comprising gabapentin, where the composition is substantially isotonic with a physiological fluid of a subject. For example, the injectable solution may be isotonic with a subject's blood or cerebrospinal fluid. Cerebrospinal fluid typically has a tonicity of about 305 mOsm. Accordingly, an embodiment of the invention provides an injectable gabapentin composition having a tonicity of about 290 mOsm to about 320 mOsm. However, such tonicities are not always achievable with gabapentin compositions. For example, gabapentin dissolved in water at a concentration of 80 mg/ml has a tonicity of about 500 mOsm. When the concentration of gabapentin in an injectable composition renders the composition hypertonic relative to a subject's physiological fluid, it is preferred that little or no amount of a tonicity enhancing agent be added to the composition. As used herein, "tonicity enhancing agent" means a compound or composition that increases tonicity of a composition. However, it will be recognized that it may be desirable to add one or more additional compounds to the composition even though the addition of the additional compound(s) will further increase tonicity of an injectable gabapentin solution. For example, it may be desirable to add to the composition an additional therapeutic agent, stabilizing compound, preservative, solubilizing agent, buffer, etc., even though tonicity will be increased.

- [33] An embodiment of the invention provides a process for preparing an injectable gabapentin composition. The method comprises mixing and/or dissolving gabapentin in a diluent or solvent to generate a composition, determining the tonicity of the composition, and adjusting the tonicity of the composition if appropriate. Any diluent or solvent may be used, provided that the diluent or solvent is pharmacologically acceptable. Preferably gabapentin is stable in the diluent or solvent. In an embodiment, water is used as the diluent. Tonicity may be determined by any means. For example, tonicity may be determined by measuring freezing point depression or decreased vapor pressure (with solutes versus substantially pure solvent). Tonicity may also be estimated. For example, a solution resulting from the mixture of one solution having a tonicity of about 500 mOsm and another solution having a tonicity of about 300 mOsm will have a resulting tonicity between about 300 mOsm and about 500 mOsm. An osmometer may be used to determine tonicity. If the tonicity of the composition is less than between about 290 mOsm to about 320 mOsm, a tonicity enhancing agent may be added to the composition to increase tonicity to between about 290 mOsm to about 320 mOsm. Any tonicity enhancing agent may be used to increase the tonicity of a composition according to various embodiments of the invention, provided that the tonicity enhancing agent is pharmacologically acceptable. Preferably, a tonicity enhancing agent is compatible with gabapentin. In an embodiment, sodium chloride is used as a tonicity enhancing agent. If the tonicity of a composition comprising gabapentin is greater than about 320 mOsm, a tonicity enhancing agent is preferably not added. In an embodiment, one or more additional agents may be added to the composition prior to determining the tonicity of the composition. For example, an additional therapeutic agent, stabilizing agent, preservative, solubilizing agent, buffer, etc. may be added prior to determining the tonicity of a composition for purposes of determining whether to add a tonicity enhancing agent.
- [34] An embodiment of the invention provides an injectable composition comprising gabapentin in a concentration of greater than about 30 mg/ml, where the injectable composition has a tonicity of less than about 900 mOsm. For example, gabapentin may be present in an injectable composition at a concentration of greater than about 31 mg/ml,

greater than about 32 mg/ml, greater than about 33 mg/ml, greater than about 34 mg/ml, greater than about 35 mg/ml, greater than about 36 mg/ml, greater than about 37 mg/ml, greater than about 38 mg/ml, greater than about 39 mg/ml, greater than about 40 mg/ml, etc., or between about 30 mg/mL to about 100 mg/mL, between about 30 mg/mL to about 90 mg/mL, between about 40 mg/mL to about 90 mg/mL, or about 80 mg/mL. An injectable gabapentin composition may have a tonicity in the range of, for example, about 250 mOsm to about 700 mOsm, in the range about 250 mOsm to about 600 mOsm, in the range of about 400 mOsm to about 550 mOsm, or about 500 mOsm.

- [35] An embodiment of the invention provides an injectable composition comprising gabapentin having a tonicity less than a corresponding composition that is the same as the injectable composition except that the corresponding composition has about 0.9% (w/v) sodium chloride. Thus, the injectable composition may comprise less than about 0.9% (w/v) sodium chloride. The composition may comprise gabapentin in any amount. Preferably, the gabapentin is present in an amount effective to treat a gabapentin-responsive disease when administered to a subject in need thereof. Gabapentin may be present in the injectable composition at a concentration of between about 0.1 mg/ml to about 100 mg/ml, between about 10 mg/ml to about 90 mg/ml, between about 20 mg/ml to about 80 mg/ml, etc. In an embodiment, gabapentin is present in an injectable composition at a concentration between about 30 mg/mL to about 100 mg/mL, between about 30 mg/mL to about 90 mg/mL, between about 40 mg/mL to about 90 mg/mL, or about 80 mg/mL. In an embodiment, a composition comprises between about 10 mg/ml and about 50 mg/ml gabapentin. For example, the composition may comprise between about 20 mg/ml and 40 mg/ml, or about 30 mg/ml.

[36] Additional Therapeutic Agents

- [37] In an embodiment, the invention provides an injectable composition comprising gabapentin and one or more additional therapeutic agents. Any additional therapeutic agent may be included in the injectable composition. Preferably, the additional therapeutic agent is compatible with gabapentin. In an embodiment, gabapentin and at

least one of the additional therapeutic agents are useful for treating the same disease state in a subject.

[38] In an embodiment, the invention provides an injectable composition comprising gabapentin and one or more additional therapeutic agents useful for treating tinnitus. Application Serial No. 10/611,459, entitled “A method for treating severe tinnitus”, filed July 1, 2003, discusses the use of intrathecally delivered gabapentin for the treatment of tinnitus. In Application Serial No. 10/611,459, local anesthetics, GABA agonists, including GABA<sub>A</sub> and GABA<sub>B</sub> agonists, serotonin agonists, thyrotropin-releasing hormones, and benzodiazapines are also discussed as being useful for treating tinnitus. Thus, according to an embodiment, the present invention provides an injectable composition comprising gabapentin and one or more of a local anesthetic, a GABA agonist, a serotonin agonist, a thyrotropin-releasing hormone, and a benzodiazapine. Specific exemplary additional therapeutic agents useful for treating tinnitus include lidocaine, bupivacaine, baclofen, muscimol, sumatriptan, sodium valproate, midazolam, alprazolam, adenosine, and pharmacologically acceptable salts thereof. Any useful amount of an additional therapeutic agent may be included in an injectable composition comprising gabapentin. Baclofen, for example, may be present in an injectable composition in a concentration between about 10 and about 4000 mcg/ml, between about 50 and about 2000 mcg/ml, between about 1000 and about 4000 mcg/ml, and between about 20 and about 2000 mcg/ml. It will be recognized that an injectable composition comprising gabapentin and one or more of the above-mentioned additional therapeutic agents may be useful for treating diseases other than tinnitus.

[39] An embodiment of the invention provides an injectable composition comprising gabapentin and one or more additional therapeutic agents useful for treatment of pain. Provisional Application Serial No. 60/513681, entitled “INTRATHECAL GABAPENTIN FOR TREATMENT OF PAIN AND EPILEPSY”, filed on October 23, 2003, which provisional application is herein incorporated by reference in its entirety, discusses the use of intrathecally delivered gabapentin for the treatment of pain and epilepsy. In Provisional Application Serial No. 60/513681 analgesics and adjuvant

analgesics are discussed as being useful for treating pain. Accordingly, an embodiment of the invention provides an injectable composition comprising gabapentin may further comprise one or more analgesic and/or adjuvant analgesic. Exemplary analgesics include opioids, NSAIDS, local anesthetics, and alpha2-adrenergic agonists. Adjuvant analgesics include anticonvulsants and antidepressants. Specific exemplary analgesics and adjuvant analgesics include, morphine, hydromorphone, bupivacaine, clonidine, baclofen, and pharmacologically acceptable salts thereof. Any useful amount of an additional therapeutic agent may be included in an injectable composition comprising gabapentin. For example, morphine sulfate may be present in an injectable composition in a concentration between about 2.5 mg/ml and about 50 mg/ml. Hydromorphone may be present in an injectable composition comprising gabapentin at, for example, a concentration of between about 1 mg/mL and about 20 mg/mL. GABA agonists, such as baclofen may also be present in a composition comprising gabapentin for treatment of pain. Any amount of a GABA agonist useful for treating pain may be used. Baclofen, for example, may be present in an injectable composition in a concentration between about 10 and about 4000 mcg/ml, between about 50 and about 2000 mcg/ml, between about 1000 and about 4000 mcg/ml, and between about 20 and about 2000 mcg/ml. It will be recognized that an injectable composition comprising gabapentin and one or more of the above-mentioned additional therapeutic agents may be useful for treating diseases other than pain.

- [40] Injectable compositions comprising gabapentin and an additional therapeutic agent according to an embodiment of the invention may be prepared in any manner that produces a product having pharmacological activity. For example, (a) an injectable composition comprising gabapentin may be mixed with an injectable composition comprising an additional therapeutic agent; (b) solid forms of gabapentin, *e.g.* gabapentin powder, and of an additional therapeutic agent may be mixed, and the resulting mixture may be added to a solvent or diluent to produce an injectable composition; (c) a solid form of gabapentin may be added to an injectable composition comprising an additional therapeutic agent; (d) a solid form of an additional therapeutic agent may be added to an injectable composition comprising gabapentin, (e) etc.

[41] In an embodiment, a sterile injectable solution comprising an additional therapeutic agent and a sterile injectable solution comprising gabapentin are added together. The sterile solutions may be added together in a sterile syringe. Adding together two sterile solutions provides for a convenient and easy means for preparing an injectable composition comprising gabapentin and an additional therapeutic agent, as well as minimizes risks of contamination associated with compounding from a non-sterile powder.

[42] Provided below in Table 1 are examples of how a sterile injectable solution comprising 80 mg/ml gabapentin may be mixed with a sterile injectable solution comprising an opioid agonist. The opioid agonists listed in Table 1 are commercially available in sterile injectable solutions. INFUMORPH is a preservative-free morphine sulfate sterile solution, and DILAUDID HP is a sterile solution comprising hydromorphone hydrochloride. Both INFUMORPH and DILAUDID HP have a tonicity of about 300 mOsm. In the examples provided in Table 1, the sterile injectable solution comprising 80 mg/ml has a tonicity of about 500 mOsm, although it will be recognized that any injectable gabapentin composition according to various embodiments of the invention may be used. A sterile injectable solution comprising 80 mg/ml gabapentin may be obtained by dissolving an appropriate amount of non-sterile gabapentin powder in water, adjusting the pH to about 6 with 1N NaOH or 1N HCl, and sterilizing the resulting solution. As shown in Table 1, sterile injectable end products may be achieved with reasonable concentrations of gabapentin and opioid. In addition, the tonicity of the resulting end products described in Table 1 are in the range of between about 300 mOsm and about 500 mOsm.

[43] Table 1: Examples of sterile injectable compositions

Common Mixtures Contemplated (V:V)	Infumorph (25 mg/mL) Final Opioid Conc (mg/ml)	Gabapentin (80 mg/mL) Final Gabapentin Conc (mg/ml)
50:50	12.5	40
90:10	22.5	8

75:25	18.75	20
10:90	2.5	72
	Dilaudid HP (10 mg/mL)	Gabapentin(80 mg/mL)
90:10	9	8
50:50	5	40
10:90	1	72

[44] It will be recognized that an injectable composition comprising gabapentin and little or no additional osmolutes may be serve as a desirable diluent for powder forms of additional therapeutic agents. Because such an injectabele gabapentin composition comprises little or no nonanalgesic osmolutes, more of an additional therapeutic agent may be added to the injectable composition while having minimal impact on the hypertonicity of the final solution. For example, with regard to tonicity, a greater amount of powdered opioid may be added to an injectable solution comprising 80 mg/ml of gabapentin in water than to a solution comprising 80 mg/ml of gabapentin in 0.9% sodium chloride.

[45] Administration

[46] Injectable compositions according to the invention may be administered to a subject through any acceptable route. For example, the compositions may be administered intravenously, subcutaneously, intrathecally, epidurally, intraparenchymally, intraperitoneally, intracerebroventricularly, etc., by infusion or injection.

[47] In an embodiment of the invention, an injectable composition comprising gabapentin is adapted for intrathecal administration. Intrathecal administration of gabapentin provides a means for achieving effective spinal concentrations of gabapentin by bypassing the saturable L-amino acid active transport system and blood-brain barrier, while reducing concomitant systemic or supraspinal drug levels. Any effective amount of gabapentin may be administered intrathecally. For example, gabapentin may be administered intrathecally in a daily dose of between about 0.1 mg and about 200 mg, between about 1 mg and about 150 mg, between about 2 mg and about 60 mg, or greater than about 25

mg. In an embodiment, gabapentin is administered in a daily dose of less than about 25 mg. For example, gabapentin may be administered at a daily dose of between about 0.1 mg and about 10 mg, between about 0.1 mg and 5 mg, between about 0.1 mg and 2 mg, between about 0.1 and 1 mg, between about 0.1 and 0.5 mg, or about 0.2 mg. It will be understood that daily dose requirements may be adjusted to account for variability in CSF volume, CSF production rates, and rate of clearance of gabapentin from the CSF. One of skill in the art will understand that such variability may be due in part to, *e.g.*, gender and/or age.

[48] Pump System

[49] Gabapentin may be administered to a subject using a therapy delivery system 15, as shown in Figure 1. The system comprises a therapy delivery device 30. The device 30 comprises a pump 40 coupled to a reservoir 12 for housing a composition comprising a therapeutic agent, such as gabapentin. The system 15 further comprises a catheter 38. The catheter 38 comprises a proximal end 35 coupled to the pump 40 and a distal end 39 adapted for delivering the composition to a desired location of a subject, *e.g.*, a patient's cerebrospinal fluid or brain tissue. It will be recognized that the catheter 38 may have one or more drug delivery regions along the length of the catheter 38 and that a drug delivery region may or may not be at the distal end 39 of the catheter 38. The therapy delivery device 30 may be implantable or may be an external device. The therapy delivery device 30 may have a port 34 into which a hypodermic needle can be inserted to inject a quantity of therapeutic agent into reservoir 12. The device 30 may have a catheter port 37, to which the proximal end 35 of catheter 38 may be coupled. The catheter port 37 may be coupled to pump 40 through an internal catheter 10. A connector 14 may be used to couple the catheter 38 to the catheter port 37 of the device 30. Device 30 may take the form of the device shown in U.S. Pat. No. 4,692,147 (Duggan), assigned to Medtronic, Inc., Minneapolis, Minn., commercially available as the Synchronmed® infusion pump, which is incorporated by reference. A system 15 comprising (a) a therapy delivery device 30 having a reservoir 12 and (b) a composition comprising gabapentin housed in the reservoir 12 is contemplated by the invention.



- [50] Therapy delivery device 30, such as Medtronic's SYNCHROMED pump system, may be operated to discharge a predetermined dosage of the pumped fluid into a subject. The therapy delivery device 30 may contain a microprocessor 42 or similar device that can be programmed to control the amount of fluid delivery. The programming may be accomplished with an external programmer/control unit via telemetry. A controlled amount of fluid comprising therapeutics may be delivered over a specified time period. With the use of a delivery device 30, different dosage regimens may be programmed for a particular patient. Additionally, different therapeutic dosages can be programmed for different combinations of fluid comprising therapeutics. Those skilled in the art will recognize that a programmed therapy delivery device 30 allows for starting conservatively with lower doses and adjusting to a more aggressive dosing scheme, if warranted, based on safety and efficacy factors.
- [51] If it is desirable to administer more than one therapeutic agent, the composition within the reservoir 12 may contain a second, third, fourth, etc. therapeutic agent. Alternatively, the therapy delivery device 30 may have more than one reservoir 12 for housing additional compositions comprising a therapeutic agent. When the device 30 has more than one reservoir 12, the pump 40 may draw fluid from the one or more reservoirs 12 and deliver the drawn fluid to the catheter 38. The device 30 may contain a valve coupled to the pump 40 for selecting from which reservoir(s) 12 to draw fluid. Further, one or more catheters 38 may be coupled to the device 30. Each catheter 38 may be adapted for delivering a therapeutic agent from one or more reservoirs 12 of the device 30. A catheter 38 may have more than one lumen. Each lumen may be adapted to deliver a therapeutic agent from one or more reservoirs 12 of the pump 40. It will also be understood that more than one implantable device 30 may be used if it is desirable to deliver more than one therapeutic agent. Such therapy delivery devices, catheters, and systems include those described in, for example, copending application Serial No. 10/245,963, entitled IMPLANTABLE DRUG DELIVERY SYSTEMS AND METHODS, filed on December 23, 2003, which application is hereby incorporated herein by reference.

- [52] A therapy delivery system 15 may be used to infuse an injectable composition in any known manner, including intravenously, subcutaneously, intrathecally, epidurally, intraparenchymally, intraperitoneally, intracerebroventricularly.
- [53] Referring to Figures 2, 3, and 4, a system or device 30 may be implanted below the skin of a patient. Preferably, the device 30 is implanted in a location where the implantation interferes as little as practicable with patient activity. Device 30 may be implanted subcutaneously in any medically acceptable area of the human body such as in a subcutaneous pocket located in the chest below the clavicle, in an abdominal subcutaneous pocket, and the like.
- [54] According to an embodiment of the invention, distal end 39 of catheter 38 is positioned to infuse a fluid into a target area of cerebrospinal fluid (CSF) of a patient. As shown in Figure 2, catheter 38 may be positioned so that the distal tip 39 of catheter 38 is located in the subarachnoid space 3 of the spinal cord between the fifth lumbar and fifth thoracic vertebrae. It will be understood that the distal tip 39 can be placed in a multitude of locations to deliver a therapeutic agent into the cerebrospinal fluid 6 of the patient. Within the spinal cord, the distal tip 39 of the catheter 38 may be inserted, for example, in the subarachnoid space 3 between the fifth thoracic (T5) and the first cervical vertebrae (C1), in the subarachnoid space 3 between the fifth lumbar (L5) and fifth thoracic vertebrae (T5), etc. The location of the distal tip 39 of the catheter 38 may be adjusted to improve therapeutic efficacy. Administering a composition comprising gabapentin at a level in the spinal canal nearer the brain may result in increased concentrations of gabapentin in the brain. Alternatively, a composition comprising gabapentin may be administered directly into the cerebral ventricles. While device 30 is shown in Figure 2, delivery of a composition comprising gabapentin into the CSF to treat epilepsy can be accomplished by injecting the therapeutic agent via port 34 to catheter 38.
- [55] Referring to FIG. 3, a system for intraparenchymal or intracerebroventricular administration of a composition comprising gabapentin is shown. Device 30 and delivery system 15 may take the form of a device and system described in US Patent No.

6,042,579, entitled "Techniques for treating neurodegenerative disorders by infusion of nerve growth factors into the brain", which patent is incorporated herein by reference in its entirety. As shown in Figure 3, the distal end of catheter 38 may terminate in a cylindrical hollow tube 38A having a distal end 115 implanted into a portion of the brain by conventional stereotactic surgical techniques. The distal portion 115 may be implanted in the brain in any medically acceptable region. In an embodiment of the invention, the distal portion 115 is implanted in a region within or proximate to an epileptic focus. In an embodiment, portion 115 comprises details as described in U.S. application Ser. No. 08/430,960, now abandoned, entitled "Intraparenchymal Infusion Catheter System," filed Apr. 28, 1995 in the name of Dennis Elsberry et al. and assigned to the same assignee as the present application, which application is herein incorporated by reference. Tube 38A may be surgically implanted through a hole in the skull 123 and catheter 38 may be implanted subcutaneously between the skull and the scalp 125 as shown in Figure 3. Catheter 38 may be joined to implanted device 30 in the manner shown and may be secured to device 30 by, for example, securing catheter 38 to catheter port 37. In an embodiment, distal end 115 of cylindrical hollow tube 38A may be implanted in a ventricle of the brain. Alternatively, the distal tip may be located in the subdural area (SD) beneath the dura under the skull 123 but outside the brain B, and within the arachnoidal space. Catheter 38 may be divided into twin tubes 38A and 38B (not shown) that are implanted into the brain bilaterally. Alternatively, tube 38B (not shown) implanted on the other side of the brain may be supplied with drugs from a separate catheter 38 and device.

- [56] As shown in Figure 4, a system for delivering therapeutic agent may include a patient-controlled activator 90, PCA. A PCA 90 may communicate with an implantable pump 40 to adjust the amount of therapeutic agent delivered. Communication between PCA 90 and implantable device 30 may be through any suitable means. In an embodiment, communication is through telemetry. Communication may be unidirectional; *i.e.*, from PCA 90 to device 30, or bi-directional. PCA 90 may be a hand held device. PCA may contain a button 92 or other suitable means for a patient to indicate a desire to alter amount of therapeutic agent delivered. Typically, a patient will depress button 92 or

activate other suitable means to direct device 30 to deliver additional therapeutic agent, such as a composition comprising gabapentin. Generally, a pulse or short-term increase in infusion rate of therapeutic agent will result as a result of the patient depressing the button 90. In an embodiment, a patient may place PCA 90 over skin in an area where device 30 is implanted. The amount and frequency of patient-controlled therapy administration may be limited by a physician or other health care provider by specifically programming the PCA 90 for a particular patient. Preferably, such programming controls would be inaccessible to the patient. It will be appreciated that a similar PCA 90 feature can be included in an external pump without the requirement of an additional device component. It will be further appreciated that while Figure 4 depicts intrathecal administration, a PCA 90 may be used with intracerebroventricular, intraparenchymal, and other routes of administration in accordance with various embodiments of the invention.

- [57] Referring to Figure 5, a system having an external therapy delivery device 30 is shown. The proximal end 35 of a catheter 38 may be coupled to the device and the distal end 39 of the catheter 39 may be positioned to deliver a therapeutic agent pumped from the external device 30 through the catheter 38 to a desired location in a subject, such as a patient's cerebral spinal fluid or brain tissue. As shown in Figure 5, the therapeutic agent, such as gabapentin, may be administered intrathecally. External delivery device 30 may be used as part of a drug trial system prior to use of an implantable pump system, examples of which are shown in Figures 2 -4. Use of an external delivery device 30 in such a manner provides an indication as to whether a patient will respond favorably to treatment prior to subjecting the patient to surgery associated with an implantable pump system. With a drug trial system, a catheter 38 may be placed to deliver a composition comprising a therapeutic agent epidurally to the patient. It will be recognized that the therapeutic agent may be administered directly to a patient's CSF as discussed above. As with the implantable delivery devices (see Figures 2- 4 and accompanying discussion), the placement position of the catheter may be varied from patient to patient or within a patient to optimize therapeutic efficacy. Any dose of therapeutic agent may be administered with an external therapy delivery device according to various embodiments

of the invention. When used as a drug trial system, the dose of a therapeutic agent is typically started conservatively with lower doses and adjusted to higher doses until pain relief is noticed. It will also be recognized that single or multiple injections, without the use of a device 30, may also be used as to screen patients that are favorable candidates for an implantable therapy delivery device.

**[58]** Kit With Instructions

**[59]** An embodiment of the invention provides a kit comprising an injectable gabapentin composition and instructions indicating that the injectable composition comprising gabapentin may be administered to cerebrospinal fluid of a subject. The instructions may include directions for administering the injectable composition comprising gabapentin to a subject's cerebrospinal fluid through any acceptable route, including for example intrathecally, intracerebroventricularly, etc or combinations thereof. The instructions may further indicate that the injectable gabapentin composition may be placed in an implantable pump 30.

**[60]** The following patent applications are generally relevant to injectable gabapentin and its use:

**[61]** US Patent Application Serial No. \_\_\_\_\_, entitled INTRATHECAL GABAPENTIN FOR TREATMENT OF PAIN, filed on even date herewith, and having Attorney Docket No. P-20216.00;

**[62]** US Patent Application Serial No. \_\_\_\_\_, entitled INJECTABLE GABAPENTIN COMPOSITIONS, filed on even date herewith, and having Attorney Docket No. P-20904.00;

**[63]** US Patent Application Serial No. \_\_\_\_\_, entitled PROCESS FOR PRODUCING INJECTABLE GABAPENTIN COMPOSITIONS, filed on even date herewith, and having Attorney Docket No. P-20907.00; and

**[64]** US Patent Application Serial No. \_\_\_\_\_, entitled INTRATHECAL GABAPENTIN FOR TREATMENT OF EPILEPSY, filed on even date herewith, and having Attorney Docket No. P-20905.00.

**[65]** All patents, patent applications, technical papers, and other publications cited herein are hereby incorporated by reference herein, each in its respective entirety. As those of ordinary skill in the art will readily appreciate upon reading the description herein, at least some of the compositions, devices and methods disclosed in the patents and publications cited herein may be modified advantageously in accordance with the teachings of the present invention.